

# Fast simultaneous plasmid vector linearization and dephosphorylation protocol

1. Prepare the following reaction mixture containing:

<b>Plasmid DNA</b>	1 µg
<b>10X Thermo Scientific FastDigest Buffer</b>	2 µL
<b>FastDigest Restriction Enzyme</b>	1 µL
<b>FastAP Thermosensitive Alkaline Phosphatase</b>	1 µL
<b>Water, nuclease-free</b>	to 20 µL
<b>Total volume</b>	<b>20 µL</b>

2. Mix thoroughly, spin briefly and incubate at 37°C for at least 15 min.

3. Stop reactions by heating at 65°C for 15 min or at 80°C for 20 min (if restriction enzyme is not inactivated at 65°C)